

PATENT Docket No. 17282CIP(AP)

AND TRADEMARK OFFICE

Examiner: Nolan, P.

Group Art Unit: 1644 RECEIVE

DEC 2 0 2002

TECH CENTER 1600/2900

'In Re Application of Steward et al

Serial No: 09/548,409; Conf. No.: 7255

Filed: April 13, 2000

For: COMPOSITIONS AND METHODS FOR

THE TREATMENT OF PANCREATITIS

TRANSMITTAL SHEET

Assistant Commissioner for Patents Washington, D.C. 20231

Sir:

Transmitted herewith is an Amendment in the above-identified application. Enclosed are:

- 1) Reply and Amendment (6 pages)
- 2) Transmittal Sheet/Certificate of Mailing/Extension of Time (2 months)
- 3) Return/Stamped Postcard

CERTIFICATE OF MAILING

I hereby certify that this correspondence is being deposited with the United States Postal Service as FIRST CLASS MAIL in an envelope addressed to: Box Amendment-Fee, Assistant Commissioner for Patents, Washington, D.C. 20231 on

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Date of Signature

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12/19/2002 DENMARU1 00000033 010885 09548409

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The fee has been calculated as shown below:

CLAIMS AS FILED

	CLAIMS REMAINING AFTER AMENDMENT		HIGHEST NO. PREVIOUSLY PAID FOR		PRESENT EXTRA	RATE	ADDITIONAL FEE
Total Claims	12	MINUS	20		= 0 ×	\$18	= \$0.00
Independent Claims	1	MINUS	3		= 0 ×	\$84	= \$0.00
If application has bee dependent claim(s),		tain multiple				\$280	= \$0.00
(Select only one)				on	e month	\$110	= \$
Time Extension Fees	:			tw	o months	\$400	= \$400.00
				th	ree months	\$920	= \$
				fοι	ur months	\$1,440	= \$*
				_	TAL ADDITION OR THIS AMENI		\$ 400.00

- () A check in the amount of \$* is enclosed (place fee in here i.e., petition, excess claims, etc.)
- (x) The Commissioner is hereby authorized to charge fees under 37 CFR 1.16 and 1.17 (associated with petition fees or excess claim fees) which may be required, or credit any overpayment to Deposit Account No. 01-0885. A duplicate copy of this sheet is enclosed.

Respectfully Submitted,

Date: 12(11(02

Signature:

Carlos A. Fisher Registration No. 36,510 Legal Department, T2-7H ALLERGAN, INC.

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

		RECEIVED
Applicant: Steward et al.	Group Art Unit: 1644	DEC 2 0 2002
Serial No.: 09/548,409) Conf. No.: 7255) Filed: April 13, 2000) For: Compositions and Methods) For the Treatment of Pancreatitis)	Examiner: Nolan, P.	TECH CENTER 1600/2900
AMEN	DMENT A	
Assistant Commissioner for Patents Washington, D.C. 20231		
Dear Sir:		
This communication is in reply to the C	Office Action mailed August 8, 200	02.
AME	NDMENT	
In the specification:		
Kindly amend the paragraph bridging pages 25 paragraph:	and 26 of the specification with the	ne following
CERTIFICATE O	F EXPRESS MAILING	
I hereby certify that this correspondence is being deposited with the Uni Amendment-Fee; Assistant Commissioner for Patents, Washington, D.C		nvelope addressed to: Box
Date of Deposit: /2/11/2002		
Person making Deposit: PONNIE FER	euson nem	
Signature: Donnie Huge	um	
D. CO. 1 /2 /1/ 2002		

Docket: 17282 CIP Serial No. 09/548,409 STEWARD et al.

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-- In a preferred embodiment, the therapeutic element is a polypeptide comprising a clostridial neurotoxin light chain or a portion thereof retaining the SNARE-protein sequence-specific endopeptidase activity of a clostridial neurotoxin light chain. The amino acid sequences of the light chain of botulinum neurotoxin (BoNT) subtypes A-G have been determined, as has the amino acid sequence of the light chain of the tetanus neurotoxin (TeNT). Each chain contains the Zn⁺⁺-binding motif **His-Glu-x-x-His** (SEQ ID: 12) (N terminal direction at the left) characteristic of Zn⁺⁺-dependent endopeptidases (HELIH in TeNT, BoNT/A /B and /E; HELNH in BoNT/C; and HELTH in BoNT/D).--

In the claims:

Kindly cancel claim 2.

Kindly amend claim 1 to read as follows:

0%) 0%) 1. (Amended) A composition for the treatment of acute pancreatitis in a mammal comprising, a first element comprising a binding element able to specifically bind a pancreatic acinar cell CCK receptor under physiological conditions, a second element comprising a translocation element derived from a clostridial neurotoxin heavy chain able to facilitate the transfer of a polypeptide across a vesicular membrane, and a third element comprising a therapeutic element derived from a clostridial neurotoxin light chain_able, when present in the cytoplasm of a pancreatic cell, to inhibit enzymatic secretion by said pancreatic cell.

REMARKS

The Examiner has requested that the sequences on page 26 of the specification be amended to incorporate a SEQ ID NO in order to relate these sequences to the paper copy of the sequence listing. Applicants have hereby made such textual reference; SEQ ID NO: 12 is a